PATENT COOPERATION TREATY



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INTERNATIONAL PRELIMINARY EXAMINATION REPORT

Anslation internal	PATENT COOPERAT		ATY	PCT/FR2003/0		
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INTERNATIONAL PRELIMINARY EXAMINATION REPORT						
	(PCT Article 36 an	d Rule 70)				
Applicant's or agent's file reference 344915D20427	FOR FURTHER ACTION			ansmittal of Internation port (Form PCT/IPEA/416		
International application No. PCT/FR2003/001873	International filing date (days 18 juin 2003 (18.06		,	day/month/year) . 2002 (18.06.2002)		
International Patent Classification (IPC) or C07D 401/12, A61K 31/506, A						
Applicant	PIERRE FABRE MED	ICAMENT				
This international preliminary examination report has been prepared by this International Preliminary Examining Authority and is transmitted to the applicant according to Article 36.						
2. This REPORT consists of a total	of sheets, includ	ing this cover sh	ieet.			
This report is also accompanied by ANNEXES, i.e., sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications made before this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions under the PCT).						
These annexes consist of a	total of sheets.					
3. This report contains indications re	elating to the following items:					
I Basis of the repor	t					
II Priority						
III Non-establishment of opinion with regard to novelty, inventive step and industrial applicability				l applicability		
IV Lack of unity of invention						
Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement						
VI Certain documents cited						
	the international application					
	ons on the international application	n				
Date of submission of the demand	Date	of completion of	f this report			
13 janvier 2004 (13.0)1.2004)	23 Sep	tember 2004	4 (23.09.2004)		
Name and mailing address of the IPEA/E	P Autho	orized officer				
Facsimile No.	Teler	hone No.				

International application No.

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Ľ	1. Basis of the report						
1.	With	regard t	o the elements of the international application:*				
	\boxtimes	the inte	emational application as originally filed				
	\boxtimes	the des	scription:				
		pages	1-22	, as originally filed			
		pages		, filed with the demand			
		pages	, filed with the letter of	, mod with the definance			
	\square	the clai					
	ΚŻ						
		pages pages		, as originally filed			
		pages	, as amended (together with any				
		pages		, filed with the demand			
	_	pages	, filed with the letter of				
		the dra	wings:	!			
		pages		, as originally filed			
		pages					
		pages	, filed with the letter of				
	$\prod t$	he seque	ence listing part of the description:				
		pages	-				
		pages					
		pages	, filed with the letter of				
_							
2.	With regard to the language, all the elements marked above were available or furnished to this Authority in the language in which the international application was filed, unless otherwise indicated under this item. These elements were available or furnished to this Authority in the following language which is: the language of a translation furnished for the purposes of international search (under Rule 23.1(b)).						
	Ш	the language of publication of the international application (under Rule 48.3(b)).					
			guage of the translation furnished for the purposes of international preliminary examination	on (under Rule 55.2 and/			
3.	With	With regard to any nucleotide and/or amino acid sequence disclosed in the international application, the international oreliminary examination was carried out on the basis of the sequence listing:					
		contain	ontained in the international application in written form.				
	\sqsubseteq		d together with the international application in computer readable form.				
			rnished subsequently to this Authority in written form.				
			nished subsequently to this Authority in computer readable form.				
		The st	The statement that the subsequently furnished written sequence listing does not go beyond the disclosure in the international application as filed has been furnished.				
		The sta	atement that the information recorded in computer readable form is identical to the writernished.	tten sequence listing has			
4.			nendments have resulted in the cancellation of:				
			the description, pages				
		- '	the claims, Nos				
			the drawings, sheets/fig				
5.		This rep beyond	oort has been established as if (some of) the amendments had not been made, since they hat the disclosure as filed, as indicated in the Supplemental Box (Rule 70.2(c)).**	ve been considered to go			
- 1	Repla in thi and 7	s report	heets which have been furnished to the receiving Office in response to an invitation under as "originally filed" and are not annexed to this report since they do not contain	Article 14 are referred to amendments (Rule 70.16			
**	Any re	eplaceme	ent sheet containing such amendments must be referred to under item 1 and annexed to this r	report.			
				-			

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IV. Lack of unity of invention					
1. In response to the invitation to restrict or pay additional fees the applicant has:					
restricted the claims.					
paid additional fees.					
paid additional fees under protest.					
neither restricted nor paid additional fees.					
2. This Authority found that the requirement of unity of invention is not complied with and chose, according to Rule 68.1, not to invite the applicant to restrict or pay additional fees.					
3. This Authority considers that the requirement of unity of invention in accordance with Rules 13.1, 13.2 and 13.3 is complied with.					
not complied with for the following reasons:					
See supplemental sheet					
,					
·					
. Consequently, the following parts of the international application were the subject of international preliminary examination in establishing this report:					
all parts.					
the parts relating to claims Nos.					

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Supplemental Box

(To be used when the space in any of the preceding boxes is not sufficient)

Continuation of: IV.

The present International Searching Authority considers that the following two subjects are not linked by a technical relationship under the terms of PCT Rule 13.2 and that, as a result, the application does not fulfil the requirement of unity of invention set forth in PCT Rule 13.1. The reasons are as follows:

Document D1 is cited in this report: D1: WO 98/22459.

In said document, example I-66 describes a compound having Formula I, wherein X=Y=CH, A=Me, B=F, D=Cl and E=F. This compound is used as a $5-HT_{1A}$ agonist. This compound has been excluded from claim 1 of the present application.

The only feature in the application that could constitute a special technical feature is Formula I. This feature is not novel (D1). Since it has not been possible to identify other features in the present application that could constitute special technical features, the present application lacks unity. The two subjects are:

- Compounds having general formula (I), wherein X=Y=CH, and the compositions and uses thereof.
- Compounds having general formula (I), wherein X=Y=N,CH, except those compounds in which X=Y=CH, and the compositions and uses thereof.

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V. Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement
Statement
Novelty (N)
Claims
1-8
YES

Claims

Inventive step (IS)

Claims

YES

Claims

1-8

NO

Industrial applicability (IA)

Claims

1-8

YES

Claims

NO

2. Citations and explanations

SUBJECT 1

The present application does not fulfil the requirements set forth in PCT Article 33(3) because the subject matter of claims 1-8 does not involve an inventive step.

Reference is made to the following document:

D1: WO 98/22459 A (KOEK WOUTER; PF MEDICAMENT (FR); BONNAUD BERNARD (FR); VACHER BERN) 28 May 1998 (1998-05-28).

The closest subject matter is described in document D1 (page 79, the table). Example I-66 describes a compound that falls within the scope of claim 1 of the present application. This compound has been excluded from claim 1 of the present application. Said compound has the same activity as the compounds of the present application.

Compound I-66 was not tested in document D1 and was only tested in the present application. Apparently, the activity of said compound is inferior to that of the compounds in example 9 (75% versus 100% stimulation, see the table on page 21).

The compounds tested in document D1 were compounds I-62, I-65 and I-55. Said compounds also have high $5-\mathrm{HT_{1A}/D2}$ selectivity.

The present application relates, in particular, to compounds having general formula (I), wherein B and E each represent a fluorine atom and D represents a chlorine atom. Compounds I-62 and I-55 also have these features but, unlike compound I-66, have not been tested with regard to stimulation in the present application.

In the present application, the problem addressed by the applicant was that of providing compounds with enhanced specific activity with respect to $5-HT_{1A}$. It is not clear that all of the compounds claimed in claim 1 have superior activity to the compounds of D1. For this reason, claims 1-8 do not involve an inventive step.

Moreover, it appears that the claim does not include all of the essential features. It was necessary to include a proviso but said proviso is not a positive feature. It is not clear why the activity of compound I-66 is inferior to that of the compounds claimed in claim 1.

SUBJECT 2

Claims 1 and 2 describe compounds having general formula (I), wherein

- (a) X or Y is CH and the other is N and they cannot be the same; and
- (b) X=Y=N.

Feature (b) of subject 2 is not supported by the description. There are no examples of feature (b) in the

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present application. It follows that feature (b) of subject 2 (claims 1, 2, 4-8) of the present application does not fulfil the requirements of PCT Article 6.

Feature (a) of subject 2 involves an inventive step. In the present application, the problem addressed by the applicant was that of providing alternative compounds having specific activity with respect to 5-HT_{1A}. The closest prior art is document D1, which describes compounds in which the piperidin-4-yl group has been substituted with a methyl pyridine group. The difference between D1 and the present application is that the compounds of the present application have a pyrimidine group, whereas the compounds of document D1 have a pyridine group. It would not have been obvious for a person skilled in the art to replace a pyridine with a pyrimidine and, as a result, feature (a) of subject 2 involves an inventive step.